

(FILE 'HOME' ENTERED AT 15:01:41 ON 26 NOV 2001)

FILE 'CAPLUS, EMBASE, BIOSIS, MEDLINE, WPIDS' ENTERED AT 15:02:11 ON 26 NOV 2001

L1           24 S (CANNE, L? OR CANNE L?)/AU,IN  
L2           16 DUP REM L1 (8 DUPLICATES REMOVED)  
L3           289 S (WILKEN, J? OR WILKEN J?)/AU,IN  
L4           4779 S (SIMON, R? OR SIMON R?)/AU,IN  
L5           1619 S (KENT, S? OR KENT S?)/AU,IN  
L6           6641 S L2 OR L3 OR L4 OR L5  
L7           13395 S (HEAD) (2A) (TAIL)  
L8           116 S (NATIVE) (2A) (CHEMICAL) (2A) (LIGAT?)  
L9           13508 S L7 OR L8  
L10          49 S L6 AND L9  
L11          2 S L10 AND (HIV OR RANTES)  
L12          2 DUP REM L11 (0 DUPLICATES REMOVED)  
L13          986916 S (DOMAIN? OR EPITOPE? OR MODULE?)  
L14          70 S (L8 OR KENT) AND L13  
L15          13 S (HETERO? OR OLIGOMER? OR FUSED OR HYBRID?) AND L14  
L16          4 DUP REM L15 (9 DUPLICATES REMOVED)  
L17          597 S L13(10A) (L7 OR L8 OR KENT)  
L18          596 S L17 AND (L7 OR L8)  
L19          0 S L18 AND CHEMOSELECT?  
L20          36 S L18 AND (HIV OR RANTES OR TRANSCRIPTION)  
L21          21 DUP REM L20 (15 DUPLICATES REMOVED)  
L22          363268 S (PROTEIN? OR PEPTIDE?) (3A) (SYNTHESES?)  
L23          71 S L22 (5A) (L8 OR L7 OR KENT)  
L24          19 S L13 AND L23  
L25          10 DUP REM L24 (9 DUPLICATES REMOVED)  
L26          506 S (CHEMICAL) (3A) (LIGAT?)  
L27          12 S L26 (10A) (FUSED OR HYBRID? OR HETERO? OR MULTIMERIC)  
L28          198 S (CROSS) (3A) (LIGAT?)  
L29          279 S (CROSS?) (3A) (LIGAT?)  
L30          8 DUP REM L27 (4 DUPLICATES REMOVED)  
L31          0 S L29 AND L8  
L32          0 S L29 AND L7  
L33          0 S L29 AND KENT  
L34          279 S L29 AND LIGAT?  
L35          0 S CHEMOSELECT? AND L29  
L36          0 S (FUSED OR FUSING OR HYBRID? OR HETERODIMER?) AND L22 AND L4  
L37          0 S L22 AND L29  
L38          24 S L26 AND DIMER?  
L39          12 DUP REM L38 (12 DUPLICATES REMOVED)

FILE 'STNGUIDE' ENTERED AT 15:37:25 ON 26 NOV 2001

FILE 'CAPLUS, EMBASE, BIOSIS, MEDLINE, WPIDS' ENTERED AT 15:37:59 ON 26 NOV 2001

FILE 'STNGUIDE' ENTERED AT 15:38:01 ON 26 NOV 2001

SET SMA OFF  
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SET SMA OFF  
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FILE 'CAPLUS, EMBASE, BIOSIS, MEDLINE, WPIDS' ENTERED AT 15:40:35 ON 26 NOV 2001

FILE 'STNGUIDE' ENTERED AT 15:40:40 ON 26 NOV 2001

FILE 'CAPLUS, EMBASE, BIOSIS, MEDLINE, WPIDS' ENTERED AT 15:41:33 ON 26

NOV 2001

FILE 'STNGUIDE' ENTERED AT 15:41:48 ON 26 NOV 2001

FILE 'CAPLUS, EMBASE, BIOSIS, MEDLINE, WPIDS' ENTERED AT 15:42:51 ON 26 NOV 2001

SET SMA OFF  
SET SMA ON  
SET SMA LOGIN

FILE 'CAPLUS' ENTERED AT 15:43:31 ON 26 NOV 2001

L41 1 S L\*\*\*

FILE 'CAPLUS, EMBASE, BIOSIS, MEDLINE, WPIDS' ENTERED AT 15:43:45 ON 26 NOV 2001

FILE 'CAPLUS, EMBASE, BIOSIS, MEDLINE, WPIDS' ENTERED AT 15:45:01 ON 26 NOV 2001

L42 0 S (PERCIPALE, P? OR PERCIPALE P?)/AU,IN

FILE 'CAPLUS, EMBASE, BIOSIS, MEDLINE, WPIDS' ENTERED AT 15:45:39 ON 26 NOV 2001

L43 31 S (PERCIPALLE, P? OR PERCIPALLE P?)/AU,IN

L44 12 DUP REM L43 (19 DUPLICATES REMOVED)

L45 208 S (CHEMICAL LIGAT?)/TI

L46 381520 S (HETERO? OR FUSED OR FUSING)/TI

L47 1128027 S (SYNTHESES?)/TI

L48 1 S L45 AND L46 AND L47

FILE 'STNGUIDE' ENTERED AT 15:48:37 ON 26 NOV 2001

FILE 'CAPLUS, EMBASE, BIOSIS, MEDLINE, WPIDS' ENTERED AT 15:49:03 ON 26 NOV 2001

L49 63706 S (DIMER? OR HETERODIMER?)/TI

L50 3 S L45 AND L49

FILE 'STNGUIDE' ENTERED AT 15:50:00 ON 26 NOV 2001

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L39 ANSWER 10 OF 12 CAPLUS COPYRIGHT 2001 ACS  
AN 1998:597749 CAPLUS  
DN 130:4078  
TI Synthesis by **chemical ligation** of homo- and heterodimeric polypeptides based on the bZIP and HTH DNA-binding motifs  
AU Guarnaccia, Corrado; Zakhарiev, Sotir; Toro, Imre; Simoncsits, Andras; Pongor, Sandor  
CS International Centre for Genetic Engineering and Biotechnology, Trieste, 34012, Italy  
SO Pept. 1996, Proc. Eur. Pept. Symp., 24th (1998), Meeting Date 1996, 441-442. Editor(s): Ramage, Robert; Epton, Roger. Publisher: Mayflower Scientific, Kingswinford, UK.  
CODEN: 66RCA5  
DT Conference  
LA English  
CC 34-4 (Amino Acids, Peptides, and Proteins)  
AB A symposium report on the prepn. of **dimeric** peptides as mimics for DNA recognition.  
ST DNA recognition peptide **dimer** mimic prepn symposium; chem ligation peptide **dimer** prepn symposium  
IT Peptides, preparation  
RL: SPN (Synthetic preparation); PREP (Preparation)  
(DNA recognition **dimer** mimics; prepn. of homo- and heterodimeric polypeptides based on the bZIP and HTH DNA-binding motifs by chem. ligation)  
IT 215722-91-1P  
RL: BPN (Biosynthetic preparation); RCT (Reactant); BIOL (Biological study); PREP (Preparation)  
(prepn. of homo- and heterodimeric polypeptides based on the bZIP and HTH DNA-binding motifs by chem. ligation)  
IT 215721-59-8P 215721-60-1P  
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)  
(prepn. of homo- and heterodimeric polypeptides based on the bZIP and HTH DNA-binding motifs by chem. ligation)  
IT 215722-47-7P 215722-50-2P 215727-80-3P  
RL: SPN (Synthetic preparation); PREP (Preparation)  
(prepn. of homo- and heterodimeric polypeptides based on the bZIP and HTH DNA-binding motifs by chem. ligation)  
RE.CNT 3  
RE  
(1) Muir, T; Biochemistry 1994, V33, P7701 CAPLUS  
(2) Percipalle, P; EMBO J 1995, V14, P3200 CAPLUS  
(3) Percipalle, P; Peptides 1994 (Proceedings of the 23rd European Peptide Symposium) 1994, P391

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L39 ANSWER 12 OF 12 CAPLUS COPYRIGHT 2001 ACS  
AN 1995:364533 CAPLUS

DUPPLICATE 8

DN 123:9921

TI **Chemical Ligation of Cysteine-Containing Peptides:**  
Synthesis of a 22 kDa Tethered **Dimer** of HIV-1 Protease

AU Baca, Manuel; Muir, Tom W.; Schnoelzer, Martina; Kent, Stephen B. H.

CS Scripps Research Institute, La Jolla, CA, 92037, USA

SO J. Am. Chem. Soc. (1995), 117(7), 1881-7

CODEN: JACSAT; ISSN: 0002-7863

DT Journal

LA English

AB Thioester-forming chemoselective reaction of unprotected peptide fragments contg. cysteine residues has been investigated. This work shows that free sulfhydryl groups are compatible with the reactive components of thioester-forming ligation chem. This allows conjugation by chem. ligation of cysteine or other thiol-contg. peptides, followed by postligation disulfide bond formation to form folded protein domains, or large multisubunit synthetic proteins. Under acidic conditions, peptides bearing bromoacetyl or .alpha.-thiocarboxylate groups did not undergo intermol. reaction with the sulfhydryl group of cysteine. Intramol. reaction also did not occur, provided a sufficient no. of intervening residues sepd. the functionalities. The results of these studies have been used in the design and synthesis of a 22 kDa tethered **dimer** HIV-1 protease analog, prep'd. by the convergent chem. ligation of four unprotected peptide segments. Two pairs of .apprx.50 residue peptides were ligated via formation of thioester bonds to form the individual monomer polypeptide chains. The ligated monomers each possessed a nonidentical two residue extension, one at the N-terminal and the other at the C-terminal, contg. an unprotected sulfhydryl group. These were subsequently linked via directed formation of a disulfide bond. The placement of the backbone thioesters and the disulfide bond were in functionally unimportant parts of the mol., and so the resulting enzyme analog retained full catalytic activity.

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